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1. (Amended) A vaccine composition for [the prophylaxis or treatment of infection in] administration to an animal, [or bird by *Lawsonia intracellularis* or related microorganism, said vaccine composition] comprising:

*a [an immunogenic,] non-pathogenic form of *L. intracellularis* or related microorganism or an immunogenic component thereof; and a pharmaceutically acceptable carrier [one or more carriers, diluents and/or adjuvants suitable for veterinary or pharmaceutical use].*

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2. (Amended) *The* vaccine composition according to Claim 1, wherein the [composition is for the prophylaxis or treatment of infection in pigs by *L. intracellularis* or related microorganism] animal is a pig.

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3. (Amended) *The* vaccine composition according to Claim 2, wherein the non-pathogenic form of *L. intracellularis* or related microorganism is an attenuated strain of the microorganism.

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4. (Amended) *The* vaccine composition according to Claim 2, wherein the non-pathogenic form of *L. intracellularis* or related microorganism is a killed preparation of the microorganism.

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5. (Amended) *The* vaccine composition according to Claim 4, wherein the [non-pathogenic form of *L. intracellularis*] killed preparation of the microorganism is a formalin-killed preparation of the microorganism.

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6. (Amended) *The* vaccine composition according to Claim 1, [or 2] wherein said [composition] immunogenic component comprises a macromolecule selected from the group

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consisting of a [peptide,] polypeptide, [protein,] a carbohydrate, a lipid [or] and a nucleic acid
[molecule or a combination thereof] from *L. intracellularis* or related microorganism, said
macromolecule being present in an amount effective to induce a protective immune response
[agent] against *L. intracellularis* or related microorganism.

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7. (Amended) *The* vaccine composition according to Claim 6, further comprising a
[wherein the composition comprises a peptide,] polypeptide[, protein or a derivative
thereof] from *L. intracellularis* or related microorganism.

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8. (Amended) A vaccine composition according to Claim 7, wherein the [peptide,]
polypeptide [or protein is in recombinant form] a recombinant polypeptide.

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9. (Amended) *The* vaccine composition according to Claim 7, further comprising a
compound selected from the group consisting of [or 8 wherein the composition comprises]
a refolding/heatshock protein, a flagellar basal body rod protein, S-adenosylmethionine:tRNA
ribosyltransferase-isomerase, autolysin, enoyl-(acyl-carrier-protein) reductase [or] and a
glucarate transporter [or derivative thereof].

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10. (Amended) *The* vaccine composition according to Claim 9, wherein the polypeptide
[protein] is GroEL having an amino acid sequence set forth in SEQ ID NO:2 or is a protein
having at least about 40% similarity thereto.

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11. (Amended) *The* vaccine composition according to Claim 9, wherein the polypeptide
[protein] is GroES having an amino acid sequence set forth in SEQ ID NO:4 or is a protein
having at least about 40% similarity thereto.

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12. (Amended) *The* vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:1 or a sequence having at least about 40% similarity thereto.

13. (Amended) *The* vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:3 or a sequence having at least about 40% similarity thereto.

14. (Amended) *The* vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:5 or a sequence having at least about 40% similarity thereto.

15. (Amended) *The* vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:6 or a sequence having at least about 40% similarity thereto.

16. (Amended) *The* vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:8 or a sequence having at least about 40% similarity thereto.

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17. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:11 or a sequence having at least about 40% similarity thereto.

18. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:13 or a sequence having at least about 40% similarity thereto.

19. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:15 or a sequence having at least about 40% similarity thereto.

20. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:17 or a sequence having at least about 40% similarity thereto.

21. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:18 or a sequence having at least about 40% similarity thereto.

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22. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:19 or a sequence having at least about 40% similarity thereto.

23. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:20 or a sequence having at least about 40% similarity thereto.

24. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:21 or a sequence having at least about 40% similarity thereto.

25. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] polypeptide [or protein] is encoded by a [nucleotide sequence] polynucleotide comprising SEQ ID NO:22 or a sequence having at least about 40% similarity thereto.

26. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] recombinant polypeptide [or protein encoded by a nucleotide sequence comprising] comprises the sequence of SEQ ID NO:7 or a sequence having at least about 40% similarity.

27. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] recombinant polypeptide [or protein encoded by a nucleotide sequence comprising] comprises the sequence of SEQ ID NO:9 or a sequence having at least about 40% similarity.

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28. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] recombinant polypeptide [or protein encoded by a nucleotide sequence comprising] comprises the sequence of SEQ ID NO:10 or a sequence having at least about 40% similarity.

29. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] recombinant polypeptide [or protein encoded by a nucleotide sequence comprising] comprises the sequence of SEQ ID NO:12 or a sequence having at least about 40% similarity.

30. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] recombinant polypeptide [or protein encoded by a nucleotide sequence comprising] comprises the sequence of SEQ ID NO:14 or a sequence having at least about 40% similarity.

31. (Amended) A vaccine composition according to Claim 8, wherein the [composition comprises a peptide,] recombinant polypeptide [or protein encoded by a nucleotide sequence comprising] comprises the sequence of SEQ ID NO:16 or a sequence having at least about 40% similarity.

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32. (Amended) A method for vaccinating an animal [or bird] against infection by *L. intracellularis* or related microorganism or treating an animal [or bird] infected by *L. intracellularis*, said method comprising the step of:

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administering to said animal [or bird] an effective amount of a non-pathogenic form of *L. intracellularis* or related microorganism or an immunogenic component thereof for a time and under conditions sufficient to induce a protective immune response against *L. intracellularis* or related microorganism.

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37. (Amended) A method according to Claim 32-[and 33] wherein said immunogenic component comprises a peptide, polypeptide, protein, carbohydrate, lipid or nucleic acid molecule or a combination thereof from *L. intracellularis* or related microorganism in an amount effective to induce a protective immune response against *L. intracellularis* or related microorganism.

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40. (Amended) A method according to [Claims 29 or 30] Claim 32, wherein the immunogenic component is a refolding/heatshock protein, a flagellar basal body rod protein, S-adenosylmethionine: tRNA ribosyltransferase-isomerase, autolysin, enoyl-(acyl-carrier-protein) reductase or a glucarate transporter or derivative thereof.

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77. (Amended) A genetic vaccine, comprising:

a polynucleotide encoding [DNA sequence having a nucleotide sequence set forth in SEQ ID NO:1 or having at least 40% similarity thereto or is capable of hybridizing to SEQ ID NO:1 under low stringency conditions, said DNA sequence capable of expression in a pig to produce an amount of] a [peptide,] polypeptide [or protein] in an amount